SEARCH REQUEST FORM

Scientific and Technical Information Center

| Requester's Full Name: Sin | J. Lee | Examiner # : 16060 Date: 11-8 | 3-10 |
|---|--|---|-------------------------|
| Art Unit: 1752 Phone | Number 30 5-0504 | Serial Number: $09/922,72$ | 3_ |
| Mail Box and Bldg/Room Location | on: <u>9805 </u> | sults Format Preferred (circle). PAPER DISK I | E-MAII |
| If more than one search is subs | mitted, please prioriti | ze searches in order of need. *********************************** | ***** |
| Include the elected species or structures, utility of the invention. Define any term known. Please attach a copy of the cover | keywords, synonyms, acro is that may have a special m r sheet, pertinent claims, and | | ept or c, if |
| Title of Invention: Positive R | esist Composition | Resist Composition | $\int_{-\infty}^{-1} t$ |
| Inventors (please provide full names): | Ogata, Tosh | riyuki; Endo, Koutaro; | |
| Earliest Priority Filing Date: | 8-7-101 | , | |
| appropriate serial number. | • | (parent, child, divisional, or issued patent numbers) along w | ith the |
| - Please search & | r a polys | siloxane resin | |
| having a hydro | xy benzyl si ise | esquioxone unit | |
| and a phenyl | Silsesyuioxane | unit sic |)3/2) |
| either — | CH 2 Si O3/2 | CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ | |
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| STAFF USE ONLY Searcher: | Type of Search NA Sequence (#) | Vendors and cost where applicable STN 78 | |
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| Date Searcher Picked Up: | Bibliographic / Can | Dr. Link | - |
| Date Completed: 11-14-02 | Litigation | Lexis/Nexis | |
| Searcher Prep & Review Time: 5 | Fulltext | Sequence Systems | - |
| Clerical Prep Time: | Patent Family | WWW/Internet | |
| Online Time: | Other | Other (specify) | - |

PTO-1590 (8-01)

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FILE 'HCAPLUS' ENTERED AT 13:52:38 ON 14 NOV 2002
          11599 S OGATA ?/AU OR TOSHIYUKI ?/AU
L1
L2
          21949 S ENDO ?/AU OR KUOTARO ?/AU
L3
            856 S KOMANO ?/AU OR HIROSHI ?/AU
L4
              5 S L1 AND L2 AND L3
                SEL L4 4 RN
     FILE 'REGISTRY' ENTERED AT 13:57:29 ON 14 NOV 2002
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              1 S L5 AND PMS/CI
     FILE 'HCAPLUS' ENTERED AT 14:01:46 ON 14 NOV 2002
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           6211 S ?SILSESOUIOXAN?
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              1 S L4 AND L7
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     FILE 'REGISTRY' ENTERED AT 14:06:55 ON 14 NOV 2002
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                STR L9
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              0 S L13 AND L14 AND L11
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     FILE 'LCA' ENTERED AT 14:21:25 ON 14 NOV 2002
L18
              1 S ?PHENYLSILSESQUIOXAN? OR (PH OR ?PHENYL?)(2A)?SILSESOUI
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              O S ?BENZYLSILSESQUIOXAN? OR (BZ OR ?BENZYL?)(2A)?SILSESQUI
     FILE 'HCAPLUS' ENTERED AT 14:24:40 ON 14 NOV 2002
L20
            819 S ?PHENYLSILSESQUIOXAN? OR (PH OR PHENYL? OR ?PHENYL) (2A)
L21
             73 S ?BENZYLSILSESQUIOXAN? OR (BZ OR ?BENZYL?) (2A) ?SILSESQUI
L22
         143167 S PHOTORESIST? OR RESIST OR RESISTS OR PHOTOMASK? OR MASK
L23
             12 S L20 AND L21
             7 S L23 AND L22
L24
L25
             48 S L21 AND L22
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L26 QUE POS OR POSITIV?
L27 26 S L25 AND L26
L28 10 S L17 OR L24
L29 5 S L23 NOT L28
L30 22 S L27 NOT (L28 OR L29)

FILE 'REGISTRY' ENTERED AT 14:32:46 ON 14 NOV 2002

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STR

L13

VAR G1=CL/O
REP G2=(0-5) C
NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
GGCAT IS UNS AT 2
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 7

STEREO ATTRIBUTES: NONE L14 STR

VAR G1=CL/O
REP G2=(0-5) C
NODE ATTRIBUTES:
CONNECT IS E1 RC AT 2
DEFAULT MLEVEL IS ATOM
GGCAT IS UNS AT 2
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 6

STEREO ATTRIBUTES: NONE

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4 ANSWERS

SEARCH TIME: 00.00.06

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L28 ANSWER 1 OF 10 HCAPLUS COPYRIGHT 2002 ACS
2002:533964 Document No. 137:101417 Mask-making using
resist having sio bond-containing polymer. Angelopoulos,
Marie; Aviram, Ari; Guarnieri, C. Richard; Huang, Wu-song; Kwong,
Ranee; Lang, Robert N.; Mahorowala, Arpan P.; Medeiros, David R.;
Moreau, Wayne M. (International Business Machines Corporation, USA).
U.S. US 6420084 B1 20020716, 9 pp. (English). CODEN: USXXAM.
APPLICATION: US 2000-602136 20000623.

AB The invention provides improved resist compns. and lithog. methods using the resist compns. which are acid-catalyzed resists characterized by the presence of an SiO-contg. polymer. The invention also encompasses methods of forming patterned material layers (esp conductive, semiconductive, or magnetic material structures) using the combination of the SiO-contg. resist and a halogen compd.-contg. pattern transfer etchant where the halogen is Cl, Br or I.

IC ICM G03F007-004

NCL 430270100

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 35, 38

ST photolithog photoresist protected polyhydroxybenzylsilsesquioxane

IT Photolithography

(UV; mask-making using resist having sio

bond-contg. polymer for)

IT Photoresists

(mask-making using resist having sio

bond-contg. polymer for)
188557-77-9DP. (4-Hydroxybenzyl)silane

IT 188557-77-9DP, (4-Hydroxybenzyl)silanetriol homopolymer, methoxycyclohexene or/and toluenesulfonate or camphorsulfonate or

adamantanecarboxylatic acid chloride protected 188629-68-7DP, methoxycyclohexene or/and toluenesulfonate or camphorsulfonate or adamantanecarboxylatic acid chloride protected

(mask-making using resist having sio bond-contg. polymer for photolithog.)

IT 24979-70-2DP, VP5000, methoxycyclohexene protected

(mask-making using resist having sio bond-contg. polymer for photolithog.)

IT 104-15-4DP, reaction product with polyhydroxybenzylsilsesquioxa ne 109-02-4DP, 4-Methylmorpholine, reaction product with polyhydroxybenzylsilsesquioxane 931-57-7DP, 1-Methoxycyclohexene, reaction product with polyvinylphenyl or polyhydroxybenzylsilsesquioxane 2094-72-6DP, 1-Adamantanecarboxylic acid chloride, reaction product with polyhydroxybenzylsilsesquioxane

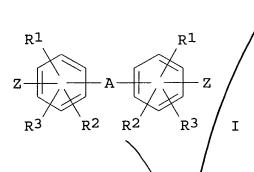
(mask-making using resist having sio bond-contg. polymer for photolithog.)

L28 ANSWER 2 OF 10 HCAPLUS COPYRIGHT 2002 ACS

2002:131262 Document No. 136:207677 Positive-working photoresist compositions and substrates equipped with photoresist layers. Ogata, Toshiyuki; Endo, Kotaro; Komano, Hiroshi (Tokyo Ohka Kogyo Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 2002055452 A2 20020220, 14 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 2000-240871 20000809.

GI

applicents



The compns contain (A) alk.-sol. polysiloxanes, (B) radiation-activated photoacid generators, and (C) compds. with their H on phenolic OH or carboxyl groups substituted with .gtoreq.1 acid dissociative groups. Preferable compds. for component (C) is given in Markush I (Z = OH, carboxyl; R1-3 = H, OH, halogen, C1-5 alkoxyl, C1-6 linear, branched, or cyclic alkyl; A = direct bond, (carboxyl-substituted) C1-5 alkylene or C2-5 alkylidene, carbonyl, Q, Q1, Q2; R4 = H, C1-5 alkyl; R5-6 = H, halogen, OH, C1-5 alkyl or alkoxy; R7-8 = C1-5 alkyl; R9-10 = H, OH, C1-5 alkyl; m = integer of 1-6) with its H on Z substituted with tertiary alkyloxycarbonylalkyl, tertiary alkyloxycarbonyl, tertiary alkyl, cyclic ether, and/or alkoxyalkyl. Substrates with a 1st resist layer consisting of an org. polymer and a 2nd 50-200 nm-thick resist layer comprising the claimed compns. are also claimed. Resist

patterns with high resoln. and excellent profiles are formed by irradn. with excimer lasers or extreme UV beams.

IT 400611-24-7P, p-Methoxybenzyltrichlorsilane-

phenyltrichlorosilane copolymer

(alk.-sol. polysiloxane-based pos. photoresist compns. contg. photoacid generators and acid-dissociative compds.)

RN 400611-24-7 HCAPLUS

CN Silane, trichloro[(4-methoxyphenyl)methyl]-, polymer with trichlorophenylsilane (9CI) (CA INDEX NAME)

CM 1

CRN 106810-48-4 CMF C8 H9 Cl3 O Si

CM 2

CRN 98-13-5 CMF C6 H5 Cl3 Si

IC ICM G03F007-039

ICS C08G077-50; C08G077-52; C08L083-06; C08L083-14; G03F007-11; H01L021-027

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 38

IT 400611-24-7P, p-Methoxybenzyltrichlorsilanephenyltrichlorosilane copolymer (alk.-sol. polysiloxane-based pos. photoresist co

(alk.-sol. polysiloxane-based pos. photoresist compns. contg. photoacid generators and acid-dissociative compds.)

L28 ANSWER 3 OF 10 HCAPLUS COPYRIGHT 2002 ACS

2001:133887 Document No. 134:185964 Radiation-sensitive resist composition. Ogata, Toshiyuki; Komano, Hiroshi (Tokyo Ohka Kogyo Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 2001051422 A2

20010223) 7 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP

AB The title compn. contains a polysiloxane and a radiation-sensitive acid generator, wherein the polysiloxane contains repeating units:

(a1) siloxane units contg. alkali sol. groups; (a2) siloxane units having acid sol. group instead of alkali sol. groups in (a1); and (a3) siloxane units having alkali-insol. groups. The compn. contg. the polysiloxane is sensitive to F2 laser and provides a pattern of the high resoln. and of the good profiles.

IT 326921-67-9DP, 4-Hydroxyphenylsilanetriol-phenylsilanetriol copolymer ester with di-tert-butyl dicarbonate, demethylated, tert-Bu carbonate esters

(silsesquioxane; radiation-sensitive resist compn.)

RN 326921-67-9 HCAPLUS

CN Silane, trichloro[(4-methoxyphenyl)methyl]-, polymer with trichlorophenylsilane, hydrolytic (9CI) (CA INDEX NAME)

CM 1

CRN 106810-48-4 CMF C8 H9 Cl3 O Si

CM 2

CRN 7732-18-5 CMF H2 O

H₂O

CM 3

CRN 98-13-5 CMF C6 H5 Cl3 Si

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Cl | Cl-si-Ph | Cl
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IC ICM G03F007-075

ICS G03F007-039; H01L021-027

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 76

IT 326921-67-9DP, 4-Hydroxyphenylsilanetriol-phenylsilanetriol copolymer ester with di-tert-butyl dicarbonate, demethylated, tert-Bu carbonate esters (silsesquioxane; radiation-sensitive resist compn.)

L28 ANSWER 4 OF 10 HCAPLUS COPYRIGHT 2002 ACS

2000:802056 Document No. 133:342499 Silicon-containing photosensitive polymers and photosensitive compositions using them. Fujiyama, Takeshi; Teramoto, Takeo (Nippon Steel Chemical Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 2000313744 A2 20001114, 6 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1999-122672 19990428.

The polymers comprise polyorganosilsesquioxanes, which are (partially) terminated with triorganosilyl groups ArORSiMe2 (Ar = o-nitrobenzyl; R = divalent org. group). The photosensitive compns. contain the polymers and photosensitizers. The compns. show high plasma resistance and give patterns with high aspect ratio. and are useful for pos. photoresists and manuf. of barrier ribs of plasma display panels.

IC ICM C08G077-388

ICS C08L083-08; G03F0Ø7-075

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 38

silsesquioxane photosensitive resist oxygen plasma resistance; pos photoresist nitrobenzyl terminated silsesquioxane; plasma display panel rib nitrobenzyl terminated silsesquioxane

IT Silsesquioxanes

(Ph) glycidyl-contg., oligomers, reaction intermediates; photosensitive nitrobenzyl-terminated silsesquioxane compns. for O plasma-resistant pos. photores; sts and PDP barrier ribs)

IT Plasma display panels

(barrier ribs; photosensitive nitrobenzyl-terminated silsesquioxane compns. for O plasma-resistant pos. photoresists and PDP barrier ribs)

IT Positive photoresists

(photosensitive nitrobenzyl-terminated silsesquioxane compns. for O plasma-resistant pos.

photoresists and PDP barrier ribs) IT Ladder polymers Silsesquioxanes (photosensitive nitrobenzyl-terminated silsesquioxane compns. for O plasma-resistant pos. photoresists and PDP barrier ribs) IT108-31-6DP, Maleic anhydride, reaction products with nitrobenzyl alc. and glycidyl-contg. Ph silsesquioxanes 126-80-7DP, reaction products with octaphenyl silsesquioxanes and nitrobenzyl-contg. carboxylic 612-25-9DP, o-Nitrobenzyl alcohol, reaction products with maleic anhydride and glycidyl-contg. Ph 51350-55-1DP, nitrobenzyl-terminated silsesquioxanes 160511-97-7DP, Phenyltrichlorosilane hydrolytic homopolymer, nitrobenzyl-terminated (photosensitive nitrobenzyl-terminated silsesquioxane compns. for O plasma-resistant pos. photoresists and PDP barrier ribs) IT 5256-79-1P, Octaphenyl silsesquioxane (reaction intermediates; photosensitive nitrobenzyl -terminated silsesquioxane compns. for O plasma-resistant pos. photoresists and PDP barrier ribs) ANSWER 5 OF 10 HCAPLUS COPYRIGHT 2002 ACS, 1999:394800 Document No. 131:80768 Electrophotographic photoreceptor having charge-transporting layer with crosslinked structure, its manufacture, and electrophotographic apparatus. Yamaguchi, Yasuhiro; Yamada, Wataru; Iwasaki, Magahiro; Nukuda, Katsumi (Fuji Xerox Co., Ltd., Japan). Jpn. Kokaj Tokkyo Koho JP 11167218 A2 19990622 Heisei, 44 pp. (Japanese). CODEN: JKXXAF. JP 1997-336195 19971205. The multilayer photoreceptor showing S-shape-type photoinduced AB potential decay has the outmost layer made of a charge-transporting layer with a crosslinked structure. The photoreceptor is manufd. by applying a soln. contg. F(LmG)n [F = n-valent org. group with charge-transporting activity; L = (hetero atom-contq.) hydrocarbylene; G = monowalent group with crosslinking reactivity; m = 0, 1; n = 1-6] and a/compd. having a group reacting with G of and processing the resulting coated layer by heat to obtain the charge-transporting layer with the crosslinked structure. The app. involves the obtained photoreceptor and a device for exposing according to digitally processed image signals. The photoreceptor

IT 228570-74-9P 2/28570-77-2P

> (multilayer electrophotog. photoreceptor involving outmost charge-transporting layer with crosslinked structure)

shows improved a rasion resistance and no change of elec. properties

RN 228570-74-9 HCAPLUS

in repeated use/.

CN Benzenepropanoic acid, 4,4'-[(3,3'-dimethyl[1,1'-biphenyl]-4,4'diyl)bis[(3,4-dimethylphenyl)imino]]bis-, bis[4-[3-(trimethoxysilyl)propyl]phenyl] ester, polymer with

triethoxyphenylsilane (9CI) (CA INDEX NAME)

CM 1

CRN 228570-73-8

CMF C72 H84 N2 O10 Si2

PAGE 1-A

$$\begin{array}{c} \text{OMe} \\ \text{OMe} \\ \text{OMe} \\ \text{OMe} \end{array}$$

PAGE 1-B

$$\begin{array}{c|c} & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & & \\ & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & &$$

CM 2

CRN 780-69-8

CMF C12 H20 O3 Si

RN 228570-77-2 HCAPLUS

CN Benzenepropanoic acid, 4-[[1,1'-biphenyl]-4-yl(3,4-dimethylphenyl)amino]-3-methyl-, 4-[3-(trimethoxysilyl)propyl]phenyl ester, polymer with trimethoxyphenylsilane (9CI) (CA INDEX NAME)

CM 1

CRN 228570-76-1 CMF C42 H47 N O5 Si

Me Me
$$CH_2-CH_2-C-O$$
 OMe CH_2) $3-Si-OMe$ OMe

CM 2

CRN 2996-92-1 CMF C9 H14 O3 Si

Ph | MeO-si-OMe | OMe

IC ICM G03G005-07

ICS G03G005-047; G03G005-05; G03G005-06

CC 74-3 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 35, 38

IT **228570-74-9P 228570-77-2P** 228570-79-4P

228570-82-9P

(multilayer electrophotog. photoreceptor involving outmost charge-transporting layer with crosslinked structure)

L28 ANSWER 6 OF 10 HCAPLUS COPYRIGHT 2002 ACS 1998:656180 Document No. 129:337644 Patterning of silsesquioxane-based

negative **photoresist** using short-wavelength light. Chokai, Minoru (Hitachi, Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 10268520 A2 19981009 Heisei, 4 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1997-74986 19970327.

AB In the process, a compn. comprising polysilsesquioxane (R1, R2 = org. group; n = p.d.) 10-90, a compd. generating strong acid under exposure of active beam or radiation light 0.01-20, and a crosslinking agent 5-60%, is applied on a substrate and irradiated with an active beam or radiation light.

IC ICM G03F007-075

ICS C08L083-04; G03F007-038; H01L021-027; H01L021-312

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 76

ST patterning silsesquioxane photoresist short wavelength light; hydroxybenzylmethylsilsesquioxane phenyliodofluoromethanesulfonic acid photoresist photolithog

IT Silsesquioxanes

(Me, hydroxycyclohexyl; silsesquioxane-based neg. photoresist with good resistance to short-wavelength light exposure)

IT Silsesquioxanes

(hydroxybenzyl, Me; silsesquioxane-based neg. photoresist with good resistance to short-wavelength light exposure)

IT Negative photoresists

Semiconductor device fabrication

(silsesquioxane-based neg. **photoresist** with good resistance to short-wavelength light exposure)

IT 80-04-6, 4,4'-Isopropylidenedicyclohexanol 556-48-9, 1,4-Cyclohexanediol

(crosslinking agent; silsesquioxane-based neg. photoresist with good resistance to short-wavelength light exposure)

IT 66003-76-7

(photoacid generator; silsesquioxane-based neg. photoresist with good resistance to short-wavelength light exposure)

L28 ANSWER 7 OF 10 HCAPLUS COPYRIGHT 2002/ACS

1996:444781 Document No. 125:234232 Photochemical reactions of silsesquioxane-based resists for ArF excimer-laser lithography. Morisawa, Taku; Fukuda, Hiroshi; Shiraishi, Hiroshi (Central Res. Ltd., Hitachi Ltd., Kokubunji, 185, Japan). Journal of Photopolymer Science and Technology, 9(3), 533-540 (English) 1996. CODEN: JSTEEW. ISSN: 0914-9244. Publisher: Technical Association of Photopolymers, Japan.

AB Two silsesquioxane polymers were investigated as ArF excimer-laser resist materials. Polyphenylmethyl silsesquioxane (PMSQ) showed pos. tone characteristics when—an—aq. base developer was

used, while polyhydroxylbenzyl silsesquioxane



(HSQ) showed neg. tone characteristics. IR spectra and x-ray photoelectron spectrometry (XPS) indicate that a major imaging mechanisms is the polarity change in PMSQ. Both materials showed sub-0.2-.mu.m resoln. capability for ArF excimer laser exposure with aq. base development.

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST silsesquioxane based photoresist excimer laser lithog; photoreaction polyphenylmethyl polyhydroxylbenzyl silsesquioxane lithog photoresist

IT Oxidation, photochemical

(in photochem. reactions of polyphenylmethyl silsesquioxane resist for ArF excimer-laser lithog.)

IT Photolysis

(photochem. reactions of polyphenylmethyl silsesquioxane resist for ArF excimer-laser lithog.)

IT Silsesquioxanes

(Me Ph, photochem. reactions of silsesquioxane-based resists for ArF excimer-laser lithog.)

IT Silsesquioxanes

(hydroxybenzyl, photochem. reactions of silsesquioxane-based resists for ArF excimer-laser lithog.)

IT Resists

(photo-, photochem. reactions of silsesquioxane-based resists for ArF excimer-laser lithog.)

IT 181429-09-4 181887-36-5

(photochem. reactions of silsesquioxane-based resists for ArF excimer-laser lithog.)

L28 ANSWER 8 OF 10 HCAPLUS COPYRIGHT 2002 ACS

1996:294909 Document No. 125:22307 Positive-working silicone resist materials. Tanaka, Haruyori; Kawai, Yoshio; Nakamura, Jiro; Matsuda, Korehito (Nippon Telegraph & Telephone, Japan). Jpn. Kokai Tokkyo Koho JP 08029987 A2 19960202 Heisei, 7 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1994-187885 19940719.

GI

OCO₂CMe₃
OH

OCO₂CMe₃
OH

(CH₂) n

(CH₂) n

(SiO₃/₂) p

(SiO₃/₂) q

I

- The title materials, which are alkali-developable and high energy ray-sensitive, contain a silicone polymer 1 (p + q = 1, x .noteq. 0; n = 1-3), an onium salt, and a N-contg. compd. The materials show high photosensitivity, high resoln., and good processability, and O plasma etching resistance. Thus, a resist comprised tert-butylcarbonated poly(hydroxybenzylsilsesquioxane), bis(tert-butylphenyl) iodonium trifluoromethanesulfonate, and o-aminobenzoic acid.
- IC ICM G03F007-075

ICS G03F007-075; G03F007-004; G03F007-029; G03F007-26; H01L021-027

- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- ST onium salt silicone resist material; nitrogen compd silsesquioxane resist material
- IT Resists

(pos.-working **resist** compn. contg. silsesquioxane and onium salt and nitrogen compd.)

IT Silsesquioxanes

(pos.-working **resist** compn. contg. silsesquioxane and onium salt and nitrogen compd.)

IT 62-53-3, Aniline, uses 95-54-5, o-Phenylenediamine, uses 99-05-8 106-50-3, p-Phenylenediamine, uses 108-45-2, 1,3-Benzenediamine, uses 118-92-3, o-Aminobenzoic acid 122-39-4, Diphenylamine, uses 150-13-0, p-Aminobenzoic acid 872-50-4, N-Methylpyrrolidone, uses 157089-26-4 157959-61-0, Bis(tert-butylphenyl)iodonium trifluoromethanesulfonate

(pos.-working **resist** compn. contg. silsesquioxane and onium salt and nitrogen compd.)

IT 158360-73-7DP, hydrolyzed, tert-Bu carbonate ester 158360-74-8DP, hydrolyzed, tert-Bu carbonate ester 158360-76-0DP, hydrolyzed, tert-Bu carbonate ester 158445-31-9DP, hydrolyzed, tert-Bu carbonate ester 158445-32-0DP, hydrolyzed, tert-Bu carbonate ester 158445-33-1DP, hydrolyzed, tert-Bu carbonate ester

(pos.-working **resist** compn. contg. silsesquioxane and onium salt and nitrogen compd.)

- L28 ANSWER 9 OF 10 HCAPLUS COPYRIGHT 2002 ACS
- 1995:226826 Document No. 122:20528 Positive-type photosensitive compositions. Aoso, Toshiaki; Mizutani, Kazuyoshi (Fuji Photo Film Co Ltd, Japan). Jpn. Kokai Tokkyo Koho JP 06011837 A2 19940121 Heisei, 63 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1991-12521 19910111.

The aq. alkali-developable title compns/ for lithog. plates, AB resists, etc., with good O plasma resistance comprise polysiloxanes contg. .gtoreq.1 mol% siloxane units formed by thermal cycloaddn. reaction of R1R2C:CR3C(SiX1X2X3):CR4R5, R1R2C:CR3CR4:CR5SiX1X2X3, R1R2C:CR3 \cancel{c} (SiR6X1X2):CR4R5, or R1R2C:CR3CR4:CR5SiR6X1X2 with QP1CF7:CR8R9, I, II, or QP1C.tplbond.CR9 and (B) 2-nitrobenzyl esters or sulfonate compds. or 2- or 3-alkoxybenzyl esters of sulfonate compds. In the formulas, R1-5 = H, (un) substity ted alkyl, aryl, silyl, siloxy; R6 = H, (un) substituted alkyl, aryl/R1R2C:CR3C:CR4R5, R1R2C:CR3CR4:CR5; R7-9 = H, (un) substituted alkyl, aryl, alkoxy, cyano, nitro, -P1Q, Q1, optionally contg. O, CO, CO2, O2C, CONR10, NR10CO, SO2, SO3; R10 = H, (un) substituted alkyl, aryl; R7R8 or R7P1 may be ring member; X1-3 = hydroxy or hydrolyzable group; P1-3 = direct bond, (un) substituted alkylene, arylene, O, CO, CO2, O2C, CONR10, NR10CO, SO2, SO3; Y = trivalent arom. group; Q = acid group of pKa below 12; Z1 = C(R7)(P1Q), CONHCO, CON(OH)CO, CON(P1Q)CO, CON(P1Q)C, (n + 2)-valent arom. group; n = 1-3.

IC ICM G03F007-075

ICS G03F007-004; G03FØ07-039; H01L021-027

CC 74-6 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

silsesquioxane photoresist nitrobenzyl ester; alkoxybenzyl ester silsesquioxane photoresist; lithog plate silsesquioxane

IT Silsesquioxanes

(photoresists and lithog. plates)

IT Resists

(photo-, sils\(\varphi\) squioxane-based)

IT 146227-70-5P 1/59448-35-8P

(in silsesquioxane-based photoresists and lithog. plates)

IT 541-59-3DP, Maleimide, reaction products with (trimethoxysilyl) butadiene-phenyltriethoxysilane silsesquioxane

(manuf. for photoresist and lithog. plates)

142-45-0DP, Acetylenedicarboxylic acid, reaction products with (trimethoxysilyl) butadiene-tolyltrimethoxysilane silsesquioxane 2210-24-4DP, N-Phenylacrylamide, reaction products with silsesquioxanes 21282-96-2DP, reaction products with silsesquioxanes 131290-90-9DP, reaction products with silsesquioxanes 159440-41-2DP, reaction products with acetylenedicarboxylic acid 159448-33-6DP, reaction products with

maleimide 159448-34-7DP, reaction products with (toluenesulfonyl)acrylamide

(manuf. for photoresists and lithog. plates)

IT 159519-43-4P 159519-44-5P (pos.-type **photoresists**)

IT 145706-02-1P 145706-03-2P 159448-32-5P

(silsesquioxane pos.-type photoresists contg.)

IT 80500-54-5 145706-09-8 159448-36-9 159448-37-0 (silsesquioxane pos.-type **photoresists** contg.)

L28 ANSWER 10 OF 10 HCAPLUS COPYRIGHT 2002 ACS

1994:311583 Document No. 120:311583 Electron-beam resist composition. Myabe, Masanori; Kobayashi, Masaichi; Nakayama, Toshimasa (Tokyo Ohka Kogyo Co Ltd, Japan). Jpn. Kokai Tokkyo Koho JP 05323609 A2 19931207 Heisei, 6 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1992-154128 19920522.

$$\begin{bmatrix} \text{HO} \longrightarrow \text{CH}_2 \text{SiO}_3/2 \end{bmatrix}_n \quad \begin{bmatrix} \text{SiO}_3/2 \end{bmatrix}_{\text{m}} \quad \text{I}$$

AB The title compn. comprises (a) alkali-sol: ladder silicone polymer having a general structure I [0.6 .ltoreq. n/(m+n) .ltoreq.0.9], (b) alkoxymethyl melamine resin, and (c) tris(dibromopropylene) isocyanurate. The compn. is useful in multilayer process of semiconductor device manuf.

IC ICM G03F007-038

ICS G03F007-075; G03F007-26; H01L021-027

- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
 Section cross-reference(s): 76
- ST photoresist compn electron beam; semiconductor device photoresist compn

IT Semiconductor devices

(electron-beam **resist** compn. in manuf. of)

IT Silsesquioxanes

(Ph, electron-beam resist compn. contg.)

IT Resists

(electron-beam, contg. alkali-sol. silicone ladder)

IT Silsesquioxanes

(hydroxybenzyl, electron-beam resist compn.

contq.)

IT 9003-08-1 82512-59-2, Tris(dibromopropyl) isocyanurate (electron-beam resist compn. contg.)

- L29 ANSWER 1 OF 5 HCAPLUS COPYRIGHT 2002 ACS/
- 2002:226226 Document No. 137:6502 Preparation of copolymerized phenylsilsesquioxane-benzylsilsesquioxane particles. Matsuda, Atsunori; Sasaki, Teruyuki; Tanaka, Toshiaki; Tatsumisago, Masahiro; Minami, Tsutomu (Department of Applied Materials Science, Graduate School of Engineering, Osaka Prefecture University, Osaka, 599-8531, Japan). Journal of Sol-Gel Science and Technology, 23(3), 247-252 (English) 2000. CODEN: JSGTEC. ISSN: 0928-0707. Publisher: Kluwer Academic Publishers.
- AB Copolymd. phenylsilsesquioxanebenzylsilsesquioxane particles were prepd. from their corresponding organotriethoxysilanes by the sol-gel method. Transparent thick films of a few microns in thickness have been successfully prepd. on glass substrates coated with indium tin oxide (ITO) by heat-treating the copolymd. particles which had been electrophoretically deposited on the substrates. The on-set temp. for thermal sintering of the copolymd. particles decreased from 150 to 50.degree.C with increasing the benzylsilsesquioxane content. These on-set temps. for thermal sintering of the particles were found to be higher by 10 to 50.degree.C than the glass transition temps. of the particles of the corresponding compn. thermal sintering of the particles should occur due to a large decrease in viscosity of the particles at temps. higher than the glass transition temps. The decrease in the on-set temp. with compn. for thermal santering as well as in the glass transition temp. of the particles can be related with the decreases in av. mol. wt. and in distribution of the mol. wt. of the particles with an increase in the benzylsilsesquioxane content.
- CC 35-5 (Chemistry of Synthetic High Polymers)
 ST phenylsilsesquioxane benzylsilsesquioxane sol
- gel polymn thermal sintering
- L29 ANSWER 2 OF 5 HCAPLUS COPYRIGHT 2002 ACS
- 2000:736520 Document No. 134:18000 Thermal softening behavior of poly(
 phenylsilsesquioxane) and poly(benzylsi/lsesquioxane
) particles. Matsuda, Atsunori; Sasaki, Teruyuki; Hasegawa, Koichi;
 Tatsumisago, Masahiro; Minami, Tsutomu (Department of Applied
 Materials Science, Graduate School of Engineering, Osaka Prefecture
 University, Sakai-shi, Osaka, 599-8531, Japan). Journal of the
 Ceramic Society of Japan, 108(Sept.), 830-835 (Japanese) 2000.
 CODEN: JCSJEW. ISSN: 0914-5400. Publisher: Ceramic Society of
 Japan.
- The thermal softening behavior of poly(phenylsilsesquioxane) (PhSiO3/2) and poly(benzylsilsesquioxane) (BnSiO3/2) particles during heat treatment has been investigated from the results of gel permeation chromatog., thermal and structural analyses of the particles. Both PhSiO3/2 and BnSiO3/2 particles thermally softened, and the on-set temp. of thermal sintering was about 140 and 50.degree C_for PhSiO3/2 and BnSiO3/2, resp. The thermal sintering of the PhSiO3/2 and BnSiO3/2 particles was caused by the decrease of viscosity at temps. higher than the glass

transition temps. Glass transition was obsd. for BnSiO3/2 in the repeated heating runs, while for PhSiO3/2 glass transition was appreciable only in the first heating run. From 29Si NMR spectra, the development of siloxane network during heat treatment was found to be more significant in PhSiO3/2 than in BnSiO3/2. The difference in the structural evolution during heat treatment makes PhSiO3/2 particles thermosetting and BnSiO3/2 ones thermoplastic.

CC 37-3 (Plastics Manufacture and Processing)
ST thermal softening polyphenylsilsesquioxane

polybenzylsilsesquioxane particle

IT Polymer morphology

(thermal softening behavior of poly(phenylsilsesquioxane

) and poly(benzylsilsesquioxane) particles)

IT Silsesquioxanes

(thermal softening behavior of poly(phenylsilsesquioxane

) and poly(benzylsilsesquioxane) particles)

IT 51350-55-1, Poly(phenylsilsesquioxane) 124741-08-8, Phenyltriethoxysilane homopolymer 186782-30-9 213474-20-5 (thermal softening behavior of poly(phenylsilsesquioxane)

) and poly(benzylsilsesquioxane) particles)

ANSWER 3 OF 5 HCAPLUS COPYRIGHT 2002 ACS L29 Document No. 129:260966 Preparation and characterization 1998:531846 of phenyl-, benzyl-, and phenethyl-subst/ituted polysilsesquioxanes. Schneider, Duane A.; Loy, Douglas A.; Baugher, Brigitta M.; Wheeler, David R.; Assink, Roger A.; Alam, Todo M.; Saunders, Randall (Encapsulants and Foams Department, Sandia National Laboratories, Albuquerque, NM, 87185-1407, USA). Polymer Preprints (American Chemical Society, Division of Polymer Chemistry, 39(2), 513-514 CODEN: ACPPAY. ISSN: 0032-3934. Publisher: (English) 1998. American Chemical Society, Division of Polymer Chemistry. Ph, benzyl, and phenethyl-substituted triethoxysilanes and AB trimethoxy silanes polymd, under acidic and basic conditions to give sol. silsesquioxane oligomers/and polymers. No gels were obsd. mol. wts. of the materials prepd. and dried at room temp. were near 2K, but would continuously increase with heating at 100.degree. to 5-15K. The glass transition temps. (Tg) for the polymers increased as the org. group was changed in the order: phenethyl < benzyl < Ph. The Tg also increased with mol. wt. The polymers were structurally characterized by H, C-13, and Si-29 NMR. The Si-29 NMR revealed substantial contributions from partially condensed silicons even with heat treatments at 200.degree.. A fully condensed polybenzylsilsesquioxane was obtained only after heating at 200.degree. with catalytic base. The inability of these monomers to form crosslinked gels under mild conditions and the difficulty encountered in increasing the mol. wt. of the sol. oligomers appears to be related to the stearic bulk of the aryl substituents. 35-5 (Chemistry ϕ f Synthetic High Polymers) CC ST

substituent effect polysilsesquioxane prepn characterization;

phenylsilsesquioxane prepn characterization;

benzylsilsesquioxane prepn characterization;

phenethylsilsesquioxne prepn characterization;

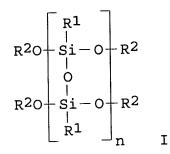
steric bulk aryl

substituent polysilsesquioxane; gel formation polysilsesquioxane substituent effect

ANSWER 4 OF 5 HCAPLUS COPYRIGHT 2002 ACS L29

Document No. 120:10471 Oxygen-resistant material for spacecraft solar cells. Sonoda, Katsumi; Kimura, Toshinori; Adachi, Hiroshi (Mitsubishi Electric Corp, Japan). Jpn. Kokai Tokkyo Koho JP 05136441 A2 19930601 Heisei, 6 pp. (Japamese). CODEN: JKXXAF. APPLICATION: JP 1991-293986 19911111.

GI



An O-resistant coating is formed to prevent surface deterioration of AB spacecraft solar cell. The O-resistant coating material may contain any material which can be used in an O atm. or an O-resistant resin. I is a silicone ladder-type resin, where R1 = Ph and/or low alkyl, R2 = H and/or low alkyl, and n = 20-100, and is the active component of the coating material.

ICM H01L031-04 IC B64G001-44; C08/005-00; C09D183-04; H01B003-46; H01L021-312; ICS H01L031-042

C08L083-00 ICI

42-10 (Coatings, Inks, and Related Products) CC Section cross-reference(s): 52

Silsesquioxanes IT

(Ph, oxygen-resistant coating contg., for spacecraft solar cell protection)

Silsesquioxanes IT

(benzyl, oxygen-resistant coating contg., for spacecraft solar cell protection),

ANSWER 5 OF 5 HCAPLUS COPYRIGHT 2002 ACS

Document No. 118:125299 / Sulfonate group-containing silsesquioxane electrolytes, and their preparation and application as solid ionic conductors.. Poinsignon, Christiane; Denoyelle, Alain; Sanchez, Jean Yves (Centre National de la Recherche Scientifique, Fr.). Fr. Demande FR 2670212 A1 19920612, 24 pp. (French). CODEN: FRXXBL. APPLICATION: FR 1990-15219 19901205. The title polyelectrolytes, useful in batteries, fuel cells, AB sensors, etc., contain the Si-bonded side groups CH2Z(SO3)t [Z =

(substituted) phenylene; t = 0-2]. Benzyltriethoxysilane was hydrolytically polymd. to give a silsesquioxane, then sulfonated to give a polymer with cond. 2 .times. 10-3 -/7 .times. 10-3 .OMEGA.-1-cm-1.

IC ICM C08G077-28

C08G077-392; H01B001-12; H01M006-18; H01M010-00; H01G009-02; ICS C08J005-20

C09D183-04 ICA

35-8 (Chemistry of Synthetic High Folymers) CC

Section cross-reference(s): 76

silsesquioxane sulfonate solid polyelectrolyte; battery ST silsesquioxane sulfonate polye/ectrolyte; fuel cell silsesquioxane sulfonate polyelectrolyte; sersor silsesquioxane sulfonate polyelectrolyte; benzyl silsesquioxane sulfonate solid polyelectrolyte

IT Silsesquioxanes

(benzyl hydrogen, sulfonated, reaction products, with divinylbenzene, prepm/. of, for polyelectrolytes)

Silsesquioxanes IT

(benzyl vinyl, di-Ph siloxane-, sulfonated, prepn. of, for solid polyelegtrolytes)

IT Silsesquioxanes

(benzyl vinyl, sulfonated, prepn. of, for solid polyelectrolytes)

Silsesquioxanes IT

IT

(benzyl, sulfonated, prepn. of, for solid

polyelectrolytes)

Siloxanes and #ilicones, compounds (di-Ph, berzyl vinyl silsesquioxane

-, sulfon ted, prepn. of, for solid polyelectrolytes)

Silsesquioxanes IT

(fluoropenzyl, sulfonated, prepn. of, for solid

polyelectrolytes)

1321-74-9DP, Divinylbenzene, reaction products with sulfonated IT hydrogen benzyl silsesquioxanes (prepn. of, for solid polyelectrolytes)

=> d 130 1-22 cbib abs hitind

ANSWER 1 OF 22 HCAPLUS COPYRIGHT 2002 ACS L30 Document No. 130:73844 Silicon polymer for chemically 1998:764063 amplified positive resist material and method of pattern formation using same. Takemura, Katsunari; Tsuchiya, Junji; Kaneko, Ichiro; Ishihara, Toshinobu (Shin-Etsu Chemical Industry Co., Ltd., Japan). Jpn. Kokai /Tokkyo Koho JP 10310642 A2 19981124 Heisei, 62 pp. (Japanese). \ ODEN: JKXXAF. APPLICATION: JP 1998-58946 19980224. PRIORITY: JP 1997-72702 19970310.

In the silicon polymer of 5,000-50,000 wt. av. mol. wt. having AB phenolic hydroxy groups, the phenolic hydrogens are partially substituted with acid unstable groups and the remaining phenolic hydroxy group are partially cross-linked inter- or/ and

- 60 66+33

intra-molecularly with -C-O-C- group. The pos. type resist material contq. the silicon polymer has a narrow range of exposure wavelength and the excellent characteristics for oxygen plasma etching. The invention provides a resist of high precision and of a high aspect ratio. IC ICM C08G077-14 ICS C08G077-46; G03F007-039; H01L021-027; G03F007-075 CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes) Section cross-reference(s): 35 silicon polymer chem amplified pos resist stIT Silsesquioxanes (Silicon polymer for chem. amplified pos. **resist** material) IT Positive photoresists (silicon polymer for chem. amplified pos/. resist material and method of pattern formation using same) IT Polysiloxanes, reactions (silicon polymer for chem. amplified pos. resist material and method of pattern formation using same) IT 109-92-2D, Ethyl vinyl ether, reaction product with poly(hydroxybenzylsilsesquioxane) 765-12-8D, Triethylene glycol divinylether, reaction product with poly(hydroxybenzylsilsesquioxane) /928-55-2D, 1-Ethoxypropene, reaction product with poly(hydroxybenzylsilsesquioxane) 999-97-3, Hexamethyldisilazane 1191-99-7, 2,3-Dihydrofuran 24424-99-5 17988-20-4 2/18148-35-7 (Silicon polymer for chem. amplified pos. resist material) IT218148-18-6 218148-20-0 218148-16-4 218148-21-1 218148-23-3 218148-26-6 218148*-*27-7 218148-28-8 218148-29-9 218148-31-3 218148-33-5 (Silicon polymer for chem. amplified pos. resist material) L30 ANSWER 2 OF 22 HCAPLUS COPYRIGHT 2002 ACS 1997:692361 Document No. 128:17350 Chemical amplification positive-working resist containing

hydroxybenzyl silsesquioxané derivative. Tsuchiya, Junji; Ishihara, Toshinobu; Nagura, Shigehiro; Takemura, Katsuya; Yamaoka, Tsugio (Shin-Etsu Chemical Industry Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 09274319 A2 19971021 Heisei, 28 CODEN: /JKXXAF. APPLICATION: JP 1996-104589 (Japanese). 19960402.

GI

=588 2844

OR OQ OH

$$(CH_2)_n \qquad (CH_2)_n \qquad (CH_2)_n$$

$$Sio_{3/2} \qquad Sio_{3/2} \qquad Z \qquad I$$

OCHMeOEt OH

$$CH_2 \qquad CH_2 \qquad CH$$

AB The title material contains (a) an org. solvent, (b) a polymer with wt. av. mol. wt. (Mw) 2000- 50,000 having a repeating unit I [R = CR1R2OR3 (R1, R2 = H, C1-6 straight chain or branched alkyl; R3 = C1-10 straight chain, branched or cyclic alkyl, R2 and R3 may link to form a ring); Q = acid-labile group; 0.05 .ltoreq. x/(x + y + z).ltoreq. 0.8, 0 .ltoreq. y/(x + y + z) .ltoreq. 0.5, 0.2 .ltoreq. z/(x + y + z) .ltoreq. 0.95; n = 1-3] as a base resin, (c) an acid-generating agent, (d) a compd. having .gtoreq.2 vinyl ether groups in its mol., and optionally (e) a compd. with mol. wt. 100-1000/having in its mol. .gtoreq.2 phenolic OH groups of which the H atoms are substituted for acid-labile groups in a ratio of 10-100 % in av., and (f) a compd. with mol. wt. 1000-3000 having in its mol. a phenolic OH group of which the H atom is substituted for the group in/a ratio of 0-60 % in av. as dissoln. inhibitor. The material shows high sensitivity toward high energy rays such as far UV rays, el'ectron beams, and x-ray and provides high resoln. patterns by development with alk. aq. solns. Thus, polymer II [x/(x + y) =0.30], (p-tert-BuOC6H4)3S+. p- MeC6H4SO3-, and [p-CH2:CHO(CH2)2OC6H4]2CMe2 were dissolved in propylene glycol monomethyl ether acetate to give a resist soln. IC

ICM G03F007-039 ICS G03F007-004; G03F007-029; G03F007-033; G03F007-075; H01L021-027

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74-5 (Radiation Chemistry, Photochemistry, and Photographic and
CC
     Other Reprographic Processes)
     hydroxybenzyl silsesquioxane photoresist
ST
     chem amplification; vinyl ether deriv pos working
     photoresist
     Silsesquioxanes
IT
        (chem. amplification resist compn. contg.
        hydroxybenzyl silsesquioxane deriv. and viny
        ether compd.)
IT
     Resists
        (pos.-working; chem. amplification resist
        compn. contg. hydroxybenzyl silsesquioxane
        deriv. and viny ether compd.)
                                                199125-57-0 199125-59-2
                                 199125-55-8
     52411-04-8
                  150610-15-4
IT
                                  199125-64-9
                    199125-63-8
     199125-61-6
         (chem. amplification resist compn. contg.
        hydroxybenzyl silsesquioxane deriv. and viny
        ether compd.)
     117458-06-7
                    123589-22-0
                                  129674-22-2
IT
        (dissoln. inhibitor; chem. amplification resist compn.
        contg. hydroxybenzyl silsesquioxane deriv.
        and viny ether compd.)
     ANSWER 3 OF 22 HCAPLUS COPYRIGHT 2002 ACS
L30
              Document No. 127:18192 Polysiloxane and positive
1997:353287
     -type resist material. Takemura, Katsuya; Tsuchiya,
     Junji; Watanabe, Osamu; Ishihara, Toshinobu (Shin-Etsu Chemical Industry Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 09087391 A2
     19970331 Heisei, 22 pp. (Japanese). CODEN: JKXXAF. APPLICATION:
     JP 1995-270580 19950925.
     The title materials comprise the silsesquioxanes
AΒ
     [SiO3/2[RQC6H4(CH2)n]]x[$iO3/2[HOC6H4(CH2)n]]y[SiO3/2[QC6H4(CH2)n]]z
     (R = alkoxy ether, furaryl, etc.; Q = \acute{acid} unstable group; x + y + z
     = 1, x, y .noteq. 0). The polymers have good alkali soly.
     polymer was prepd. by/reaction of hydroxybenzylsilanetriol
     silsesquioxane and chloromethyl\Me/ether.
     ICM C08G077-14
IC
          G03F007-004; G63F007-029; G03F007-039; G03F007-075; G03F007-30;
     ICS
          H01L021-027
                                                        =569139b
     35-8 (Chemistry of Synthetic High Polymers)
CC
     Section cross-ref/erence(s): 74
     hydroxybenzylsilanetriol silsesquioxane deriv
ST
     photoresist; pos resist silsesquioxane
     deriv
IT
     Resists
         (pos.-working; polysiloxane and pos.-type
        resist mat/erial)
     Polysiloxane/s, preparation
IT
     Silsesquioxanes
         (reaction products with ethers; polysiloxane and pos
         .-type resist material)
     107-30-2DP, Chloromethyl methyl ether, reaction products with
IT
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hydroxybenzyl silsesquioxane 109-53-5DP, Iso-butyl vinyl ether, reaction products with hydroxybenzyl 109-92-2DP, Ethyl vinyl ether, reaction silsesquioxane products with hydroxybenzyl silsesquioxane 111-34-2DP, n-Butyl vinyl ether, reaction products with hydroxybenzyl silsesquioxane 926-02-3DP, tert-Butyl vinyl ether, reaction products with hydroxybenzyl 1191-99-7DP, 2,3-Dihydrofuran, reaction silsesquioxane products with hydroxybenzyl silsesquioxane 4525-32-0DP, Dibutyl dicarbonate, reaction products with hydroxybenzyl silsesquioxane 188557-77-9DP, reaction products with ethers 188629-68-7DP, reaction products with ethers (polysiloxane and pos.-type resist material)

L30 ANSWER 4 OF 22 HCAPLUS COPYRIGHT 2002 ACS
1997:178248 Document No. 126:179065 Chemically sensitized
positive-working silsesquioxane resist material.
Takemura, Katsuya; Tsucha, Junji; Ishihara, Toshinobu (Shinetsu Chem Ind Co, Japan). Jpn. Kokai Tokkyo Koho JP 08334901 A2 19961217
Heisei, 16 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP
1996-94846 19960325. PRIORITY: JP 1995-103125 19950404.

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

The material comprises an acid generator and a polysiloxane AB (silsesquioxane) expressed by R(OSiMe3)aSiO(3-a)/2 (A; R = X1, X2, X3; Q = group unstable to an acid (excluding trimethylsily1 group); polysiloxane at least contains X1/and X2 group; m = pos./ no. satisfying 0.001 .ltoreq. m $\sqrt{1}$ toreq. 0.05; n = 1, 2; 3). polysiloxane having a repeating unit I is treated with trimethylsilyl iodide for trimethylsilylation of silanol group and of Me group in methoxy group, Mydrolyzed to generate phenolic OH group, treated with hexamethy/silazane or trimethy/silyl chloride for trimethylsilylation of a part the H atoms in the phenolic OH group and of H atoms in the residual silanol group to give a polysiloxane expressed by I/I in which a part or all of the residual phenolic OH group is substituted with a group unstable to an acid. The resist shows high sensitivity for high-energy ray such as far-UV, electron ray, /and x ray, and is developable by an alk. aq. soln. and suited for forming minute patterns for semiconductor device manuf.

IC ICM G03F007-039 / ICS G03F007-004; G03F007-075; H01L021-027

GΙ

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 37, 38, 76

ST pos working resist silsesquioxane alkali

5131126

developable; semiconductor device manuf resist silsesquioxane

IT Resists

Semiconductor devices

(prepn. of silsesquioxane for pos.-working resist)

IT Silsesquioxanes

(prepn. of silsesquioxane for pos.-working resist)

IT 157089-24-2

(acid generator in resist; prepn. of silsesquioxane for pos.-working resist)

75-77-4DP, Trimethylsilyl chloride, reaction products with trimethylsilylated and hydrolyzed methoxybenzyl silsesquioxane, tert-butoxycarbonate 999-97-3DP, Hexamethyldisilazane, reaction products with trimethylsilylated and hydrolyzed methoxybenzyl silsesquioxane, tert-butoxycarbonate 16029-98-4DP, Trimethylsilyl iodide, reaction products with methoxybenzyl silsesquioxane, hydrolyzed, trimethylsilylated, tert-butoxycarbonate 24424-99-5DP, reaction product with trimethoxysilyloxy silsesquioxane 161055-58-9DP, hydrolyzed and trimethylsilylated, tert-butoxycarbonate

(prepn. of silsesquioxane for **pos**.-working **resist**)

L30 ANSWER 5 OF 22 HCAPLUS COPYRIGHT 2002 ACS
1997:174546 Document No. 126:179064 Chemically sensitized
positive-working resist material containing
polysiloxane. Takemura, Katsuya; Tsucha, Junji; Ishihara, Toshinobu
(Shinetsu Chem Ind Co, Japan). Jpn. Kokai Tokkyo Koho JP 08334900
A2 19961217 Heisei, 13 pp. (Japanese). CODEN: JKXXAF.
APPLICATION: JP 1996-94845 19960325. PRIORITY: JP 1995-103124
19950404.

<131126

GI

$$-- (CH2)n - OQ = X1 - (CH2)n - OH = X2$$

Ι

II

- The material comprises an acid generator and a polysiloxane AB (silsesquioxane) expressed by R(OSiMe3)mSiO(3-m)/2 (A; R = X1, X2; polysiloxane contains .gtoreq.1 X1 group; Q = group unstable to an acid; m = pos. no. satisfying 0.001 .ltoreq. m .ltoreq. 0.05; n = 1, 2, 3). A polysiloxane having a repeating unit I is treated with hexamethyldisilazane or trimethylsilyl chloride to trimethylsilylation of H in silanol group, then treated with trimethylsilyl iodide to convert Me group in methoxy group to trimethylsilyl group, hydrolyzed to generate a polysiloxane having a repeating unit II and having a phenolic OH group, and then treated by partially (or completely) substituting the phenolic OH group with a group unstable to an acid to give A which is used as the resist component. The resist shows high sensitivity for high-energy ray such as far-UV, electron ray, and x ray, and is developable by an alk. aq. soln. and suited for forming minute patterns for semiconductor device manuf.
- IC ICM G03F007-039

ICS G03F007-004; G03F007-075; H01L021-027

- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
 Section cross-reference(s): 37, 38, 76
- ST pos working resist silsesquioxane alkali developable; semiconductor device manuf resist silsesquioxane
- IT Resists

(prepn. of hydroxybenzyl silsesquioxane for pos.-working resist)

IT Semiconductor devices

(prepn. of hydroxybenzyl silsesquioxane for pos.-working resist for manuf. of)

IT Silsesquioxanes

(prepn. of hydroxybenzyl silsesquioxane for pos.-working resist for manuf. of)

IT 157089-24-2

(acid generator in resist; prepn. of hydroxybenzyl silsesquioxane for pos.-working resist)

- IT 75-77-4DP, Trimethylsilyl chloride, reaction products with methoxybenzyl silsesquioxane and trimethylsilyl iodide, hydrolyzed, tert-butoxycarbonate 999-97-3DP. Hexamethyldisilazane, reaction products with methoxybenzyl silsesquioxane and trimethylsilyl iodide, hydrolyzed, tert-butoxycarbonate 16029-98-4DP, Trimethylsilyl iodide, reaction products with methoxybenzyl silsesquioxane and hexamethyldisilazane, hydrolyzed, tert-butoxycarbonate 34619-03-9DP, reaction product with hydroxybenzyl 161055-58-9DP, psilsesquioxane Methoxybenzyltrichlorosilane hydrolytic homopolymer, trimethylsilylated, hydrolyzed, tert-butoxycarbonate (prepn. of hydroxybenzyl silsesquioxane for pos.-working resist)
- L30 ANSWER 6 OF 22 HCAPLUS COPYRIGHT 2002 ACS
- 1996:534088 Document No. 125:181352 Manufacture of self-developable positively working pattern with aromatic siloxane or silsesquioxane. Tsucha, Hiroko; Shiraishi, Hiroshi; Fukuda, Hiroshi; Terasawa, Tsuneo (Hitachi Ltd, Japan; Hitachi Chemical Co Ltd). Jpn. Kokai Tokkyo Koho JP 08181132 A2 19960712 Heisei, 5 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1994-319784 19941222.
- AB The title method comprises the following processes (1) applying a resist film mainly contg. a polymer having a unit in which Si has linkages with an arom. group-contg. substituent and O and (2) irradiating an active chem. beam to vaporize the irradiated part of the resist film. The patterned resist film may be used for an etching mask of the other resist film. The resist pattern showed good dry-etching resistance.
- IC ICM H01L021-312 ICS H01L021-3065
- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- ST pos working resist pattern siloxane silsesquioxane; self developable photoresist dry etching resistance; mask etching resist pos siloxane arom
- IT Siloxanes and Silicones, preparation Silsesquioxanes

(manuf. of self-developable **pos.**-working **resist** pattern for etching **mask** by using arom. siloxane or silsesquioxane)

IT Silsesquioxanes

(hydroxybenzyl, manuf of self-developable pos.-working resist pattern for etching mask by using arom. siloxane or silsesquioxane)

IT Resists

(photo-, manuf. of self-developable **pos**.-working **resist** pattern for etching **mask** by using arom. siloxane or silpesquioxane)

IT 29226-39-9P, Diphenylsilanediol homopolymer 32129-24-1P,
Diphenylsilanediol homopolymer, sru 51350-55-1P, Phenylsilanetriol
homopolymer, ladder sru 157374-41-9P, Phenylsilanetriol
homopolymer

(manuf. of self-developable **pos**.-working **resist** pattern for etching **mask** by using arom. siloxane or silsesquioxane)

L30 ANSWER 7 OF 22 HCAPLUS COPYRIGHT 2002/ACS
1995:367480 Document No. 122:147310 Acid sensitive resist
and patterning of same. (International Business Machines Corp.,
USA). Jpn. Kokai Tokkyo Koho JP 06184311 A2 19940705 Heisei, 9 pp.
(Japanese). CODEN: JKXXAF. APPLICATION: JP 1993-210073 19930825.
PRIORITY: US 1992-943086 19920910.

GΙ

The title pos.-working resist contains acid-sensitive arylsilsesquioxanes (I; m = 0, 1; n .gtoreq. 3; .gtoreq. 15% of R is tert-butyloxycarbonyl, secondary alkyloxycarbonyl, other inactive benzyloxycarbonyl with the remainder H] contg. protected phenolic OHs in a pendant group capable of yielding phenolic OHs upon reaction with acids, the protective group serving as a dissoln. inhibitor. The title patterning is effected by coating a substrate with the resist compn. contg. the above polymer, a photo acid generator, and anthracene-type sensitizers, baking, patternwise exposing, and developing with alkali.

IC ICM C08G077-38

ICS C08L083-06; G03F007/075

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST photoresist acid generating pos working;

silsesquioxane pos working resist

IT Silsesquioxanes

(photoresist comp/ns. contg. acid-sensitive)

IT Resists

(photo-, acid-s∉nsitive silsesquioxane)

IT 24424-99-5DP, Di-tert-butyl dicarbonate, reaction product with poly(hydroxybenzylsilsesquioxane)

(pos.-working photoresist compn. contg.)

IT 161055-58-9P, Trichloro[(4-methoxyphenyl)methyl]silane hydrolytic homopolymer 161099-32-7DP, hydrolyzed, esters with di-tert-Bu dicarbonate

(pos.-working photoresist compn. from)

L30 ANSWER 8 OF 22 HCAPLUS COPYRIGHT 2002 AÇS

1994:641833 Document No. 121:241833 Positive-working
resist materials consisting of silicone resin and onium
salt. Tanaka, Haruyori; Kawai, Yoshio; Matsuda, Korehito (Nippon
Telegraph & Telephone, Japan). Jpn. Kokai Tokkyo Koho JP 06118651
A2 19940428 Heisei, 7 pp. (Japanese). CODEN: JKXXAF. APPLICATION:
JP 1992-294009 19921008.

OCO₂CMe₃
OH
$$(CH2)n$$

$$(CH2)n$$

$$(SiO3/2)x
$$(SiO3/2)m$$
I$$

The title resist materials can be developed by aq. base and are sensitive to high energy rays, and contain a silicone polymer I (x + m = 1; x .noteq. 0; n = 1-3) and RpAM [R = (substituted) arom. group; A = sulfonium, iodonium; M = MeC6H4SO3-, CF3SO3-; p = 2, 3]. These resist show good photosensitivity, high resoln., and improved processability. Thus, (ethoxybenzyl)trichlorosilane was hydrolyzed, polymd., treated with trimethylsilyl iodide, and then with di-tert-Bu dicarbonate to give poly(p-hydroxybenzylsilsesquioxane) tert-Bu carbonate (II). A typical resist contains II and (p-MeO6H4)PhI+ p-MeC6H4SO3-.

IC ICM G03F007-039

ICS G03F007-004; G03F007-029; G03F007-075; H01L021-027

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST pos working resist silicone onium

IT Silsesquioxanes

(prepn. of, resists contg. oniums and, pos.-working, alkali-developable)

IT Resists

(pos.-working, alkali-developable, contg. silicones and oniums)

158360-73-7DP, hydrolyzed, tert-Bu carbonate 158360-74-8DP, hydrolyzed, tert-Bu carbonate 158360-76-0DP, hydrolyzed, tert-Bu carbonate 158445-31-9DP, hydrolyzed, tert-Bu carbonate 158445-32-0DP, hydrolyzed, tert-Bu carbonate 158445-33-1DP, hydrolyzed, tert-Bu carbonate

(prepn. of, resists contg. oniums and, pos

.-working, alkali-developable)

IT 27126-77-8 84563-54-2 110928-18-2 111281-12-0 115298-63-0 141801-36-7 154093-57-9 156184-16-6 156184-17-7 157692-55-2 157692-56-3 158360-70-4 158360-71-5

(resists contg. silicones and, pos.-working, alkali-developable)

L30 ANSWER 9 OF 22 HCAPLUS COPYRIGHT 2002 ACS

1994:334953 Document No. 120:334953 High-sensitivity **positive**-working **photoresist** for high-resolution patterning.
Sakamizu, Toshio; Shiraishi, Hiroshi; Sugyama, Hisashi (Hitachi Ltd,

Japan; Hitachi Chemical Co Ltd). Jpn. Kokai Tokkyo Koho JP 05188597 A2 19930730 Heisei, 5 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1992-2065 19920109.

- AB The title **photoresist** compn. contains an alkali-sol. silicone, contg. phenolic-group-contg. side chains with the OH(s) substituted by acid attack-releasable groups, and a photo acid generator. The acid attack-releasable groups are .gtoreq.1 selected from tetrahydropyranyl, 2-methoxy tetrahydropyranyl, 4-methoxytetrahydropyranyl, 2-cthoxytetrahydropyranyl, 2-methoxy-4-methyltetrahydropyranyl, and tert-butoxy.
- IC ICM G03F007-039

ICS G03F007-004; G03F007-028; G03F007-075; G03F007-26; H01L021-027

- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- ST photoresist silicone tetrahydropyranyl butoxy; silsesquioxane photoresist
- IT Silsesquioxanes

(polyhydroxybenzyl-, tetrahydropyranyl- or

butoxy-substitutéd, photoresist compn.-contg..)

IT Resists

(photo-, tetrahydropyranyl- or tert-butoxy-substituted silicones-based)

- IT 110-87-2D, 3,4-Dihydro-2H-pyran, reaction product with polyhydroxybenzylsilsesquioxane 4454-05-1D, 3,4-Dihydro-2-methoxy-2H-pyran, reaction product with polyhydroxybenzylsilsesquioxane 51300-90-4D, reaction product with polyhydroxybenzylsilsesquioxane (photoresist compn. contg.)
- L30 ANSWER 10 OF 22 HCAPLUS COPYRIGHT 2002 ACS
- 1994:148960 Document No. 120:148960 Silicon-containing

 positive resist and method of using the same in
 thin film packaging technology. Sachdev, Krishna G.; Sachdev,
 Harbans S.; Whitaker, Joel R. (International Business Machines
 Corp., USA). Eur. Pat. Appl. EP 568476 A2 19931103, 12 pp.
 DESIGNATED STATES: R: DE, FR, GB. (English). CODEN: EPXXDW.
 APPLICATION: EP 1993-480025 19930323. PRIORITY: US 1992-876277
 19920430
- AB Photosensitive Si-contg. resist compns. comprises silsesquioxane and arom. siloxane esters with diazonaphthoquinone sulfonyl groups for imageable 0 RIE barrier films.
- IC ICM G03F007-075 ICS C08G077-38
- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- ST silsesquioxane pos resist thin film packaging
- IT Packaging materials

(thin film pos. regist for)

IT Silsesquioxanes

(benzyl, for thin film pos. resist)

IT Resists

(photo-, pos., silsesquioxane for)

=542223

- L30 ANSWER 11 OF 22 HCAPLUS COPYRIGHT 2002 ACS
- 1994:148593 Document No. 120:148593 Evaluation of a deep-UV bilayer resist for sub-half micron lithography. Brunsvold, W.; Stewart, K.; Jagannathan, P.; Sooriyakumaran, R.; Parrill, J.; Muller, K. P.; Sachdev, H. (Technol. Prod. Div., IBM, Hopewell Jct., NY, 12533, USA). Proceedings of SPIE-The International Society for Optical Engineering, 1925(Advances in Resist Technology and Processing X), 377-87 (English) 1993. CODEN: PSISDG. ISSN: 0277-786X.
- AB A chem. amplified silicon-contg. resist has been formulated and evaluated as a thin imaging layer in a pos. tone deep-UV (DUV) bilayer scheme. The key component is a silicon-contg. polymer which has been charagterized by GPC, UV, and dissoln. rate studies. Dose and focus latitudes were detd. for 0.4 and 0.5 mm patterns exposed on a SVGL Micrascan I step and scan system and on KrF excimer laser steppers/. The dose latitude on a GCA (0.35 NA) excimer was found to be 20% for 0.4 um features, and about 30% for 0.5 um features (.+-.10% CD variation). Focus latitude was at least 2 um for 0.5 um patterns. Wafer to wafer LW uniformity as well as within wafer uniformity will be shown. Typical processing involves 5-10 mJ/cm2 exposure doses, employing a 90 .degree. post-expose bake (PEB) and a 60s development in 0.21N tetramethylammonium hydroxide. The dependence of linewidth upon PEB was found to be about 13 nm per/degree C for 0.5 um features. Pattern transfer into the hardbaked i-line resist underlayer was done in an MLR chamber on an AME 5000. A low pressure etch is preferred to eliminate residue but this can lead to a higher non-uniformity across the wafer. Sidewall roughness was prevalent and this could be partially attributed to "feet" on the silicon-contg. imaging layer.
- CC 74-5 (Radiation Chemistry Photochemistry, and Photographic and Other Reprographic Processes)
- ST hydroxybenzylsilsesquioxane butyloxycarbonyl modified bilayer photoresist lithog; silsesquioxane butyloxycarbonyl modified bilayer photoresist lithog

 IT Silsesquioxanes

Silsesquioxanes / (lithog. chem. amplified **photoresist** layer contg., for deep-UV **pos**. tone/bilayer scheme)

IT Silsesquioxanes (hydroxybenzyl, butyloxycarbonyl-modified, as lithog.

deep-UV photoresist for bilayer systems)
Resists

(photo-, chem. amplified, butyloxycarbonyl-modified hydroxybenzyls; ilsesquioxane as top layer for bilayer system, for deep-UV lithog.)

TT 75-59-2, Tetramethylammonium hydroxide
(lithog. developer, for chem. amplified photoresist
imaging layer based on butyloxycarbonyl-modified
hydroxybenzylsilsesquioxane)

L30 ANSWER 12 OF 22 HCAPLUS COPYRIGHT 2002 ACS

IT

- 1993:682009 Document No. 119:282009 New silicon containing positive resist and its applications for sub-half micron lithography. Sachdev, H. S.; Whitaker, J. R.; Sachdev, K. G. (IBM E. Fishkill, Hopewell Junction, NY, USA). Microelectronic Engineering, 21(1-4), 223-6 (English) 1993. CODEN: MIENEF. ISSN: 0167-9317.
- New silicon contg. photosensitive polymers were synthesized using poly(p-hydroxybenzylsilsesquioxane-p-methoxybenzylsilsesquioxane) resins as starting materials and partially replacing the available (OH) groups with 2,1-diazonaphthoquinone sulfonyloxy molety. Bilayer resist formulations derived from these polymers have higher sensitivity and contrast as compared to the resists based on silicon resins/PAC mixts. for sub-micron and sub-half micron i-line, deep-UV, and g-line lithog.

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

photoresists silicone polymer microlithog;
hydroxybenzylsilsesquioxane methoxybenzylsilsesquioxane
diazonaphthoquinone sulfonyloxy polymer photoresist;
silsesquioxane diazonaphthoquinone sulfonyloxy substituted
photoresists

IT Silsesquioxanes (methoxybenzyl, partially demethylated and functionalized with diazonaphthoquinone sulfonyl, photoresist for sub-half micron lithog. from)

IT Resists

(photo-, pos.-working, hydroxybenzylsilsesquioxane -methoxybenzylsilsesquioxane polymer partially functionalized with diazonaphthoquinone sulfonyloxy as)

IT 28020-74-8D, reaction product/with partially demethylated methoxybenzylsilsesquioxane / 114747-43-2D, reaction product with partially demethylated methoxybenzylsilsesquioxane (photoresist for sub-half micron lithog. from)

L30 ANSWER 13 OF 22 HCAPLUS COPYRIGHT 2002 ACS
1992:560946 Document No. 117:160946 Positive-working
resist compositions using alkali-soluble ladder silicone
polymer. Tokutake, Nobuo; Obara, Hidekatsu; Tanaka, Hatsuyuki;
Nakayama, Toshimasa (Tokyo Ohka Kogyo K. K., Japan). Jpn. Kokai
Tokkyo Koho JP 04130324 A2 19920501 Heisei, 6 pp. (Japanese).
CODEN: JKXXAF. APPLICATION: JP 1990-250332 19900921.

The resist compns. contain a alkali-sol. ladder silicone polymer [p-HOC6H4CH2SiO3/2]n[RhCH2SiO3/2]m (I; 0.5 .ltoreq. n/(n + m) .ltoreq. 0.7) and a photosensitive compd. The compns. show good resistance to O plasma and provide high resoln. patterns. Thus, a resist compn. comprised I and a condensate of naphthoquinonediazido-5-sulfonic acid and 2,3,4-trihydroxybenzophenone.

IC ICM G03F007-075

ICS G03F007-022; G03F007-039; H01L021-027

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and

Other Reprographic Processes) Section cross-reference(s): 76 STpos working photoresist silsesquioxane IT Silsesquioxanes (hydroxybenzyl, pos.-working photoresist contg., with quinonediazide compd.) IT (photo-, pos.-working, using hydroxybenzylsilsesquioxanes, for good oxygen plasma resistance) IT 53050-67-2 (pos.-working photoresist contq., with silsesquioxanes) L30 ANSWER 14 OF 22 HCAPLUS COPYRIGHT 2002 ACS 1992:184426 Document No. 116:184426 Study on organosilicon positive resist. III. Organosilicon positive excimer laser resist (OSPR-2016). Sugiyama, Hisashi; Mizushima, Akiko; Nate, Kazuo (Prod. Eng. Res. Lab., Hitachi Ltd., Yokohama, 244, Japan). Journal of Applied Polymer Science, 44(9), 1591-4 (English) 1992. CODEN: JAPNAB. ISSN: 0021-8995. A new alkali-developable organosilicon pos. excimer laser AB (KrF) resist (OSPR-2016) has been developed for a bilayer resist system. OSPR-2016 is composed of poly(phydroxybenzylsilsesquioxane) and Me cholate-tris(.alpha.diazoacetoacetate). The ratio is 72.5:27.5 wt./wt. A sample of 0.5-.mu.m thick O&PR-2016 resolved 0.35 .mu.m L&S patterns when exposed to a dose of 320 mJ/cm2 from an excimer laser projection printer (NA = 0.37) 74-5 (Radiation Chemistry, Photochemistry, and Photographic and CC Other Reprographic Processes) ST organosilicon pos photoresist excimer laser submicron IT Silsesquioxanes (hydroxybenzyl, excimer laser pos. resist system contg., for submicron lithog.) IT Resists (photo-, pos. organosilicon, excimer-laser, for submicron lithog.) IT 631-41-4 (developer, for pos. organosilicon resist system for submicron lithog.) IT 140608-65-7, OSPR-2016 (excimer-laser pos. resist system, for submicron lithog.) IT 123153-97-9 (hydroxybenzyl, excimer laser pos. resist

L30 ANSWER 15 OF 22 HCAPLUS COPYRIGHT 2002 ACS 1992:184425 Document No. 116:184425 Study on organosilicon

system contq., for submicron lithog.)

positive resist. II. Organosilicon positive photoresist (OSPR-1334) and its application to bilayer resist system. Sugiyama, Hisashi; Mizushima, Akiko; Inoue, Takashi; Nate, Kazuo (Prod. Eng. Res. Lab., Hitachi Ltd., Yokohama, 244, Japan). Journal of Applied Polymer Science, 44(9), 1583-90 (English) 1992. CODEN: JAPNAB. 0021-8995.

- AΒ A new alkali-developable organosilicon pos. photoresist (OSPR-1334) and a bilayer resist process with OSPR-1334 has been developed. OSPR-1334 is composed of poly(p-hydroxybenzylsilsesquioxane) and naphthoquinone diazide. The sensitivity and the resoln. are almost the same as those of conventional novolak-based resists when aq. tetrakis(2-hydroxyethyl) ammonium hydroxide is used as a developer. Also, OSPR-1334 has excellent resistance to O2 RIE. The etching rate is 3.6 nm/min, whereas that of polyimide resins or hard-baked novolak-based resists is 100/nm/min. Submicron patterns with a high aspect ratio tan easily be obtained with this bilayer resist process.
- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- STorganosilicon pos photoresist bilayer submicron

IT Silsesquioxanes

(hydroxybenzyl, pos. photoresist

compn. contg., for submicron lithog.)

IT

(photo-, pos. organosilicon, for bilayer resist system for submicron lithog.)

- IT631-41-4, Tetrakis (2-hydroxyethyl) ammonium hydroxide (developer, for organosilicon pos. resist for submicron lithog.)
- IT 5610-94-6

(organosilicon pos. resist contg. photoactive compd. from)

ΙŢ 120366-99-6, OSPR-1334

> (submicron lithog bilayer resist system contga pos.)

L30 ANSWER 16 OF 22 HCAPLUS COPYRIGHT 2002 ACS

Document No. 116:152548 Study on organosilicon positive resist. Syntheses and I. characterization of silsesquioxane, sixoxane, and silmethylene polymers with phenolic hydroxy groups. Sugiyama, Hisashi; Inoue, Takashi; Nate, Kazuo (Prod. Eng. Res. Lab., Hitachi Ltd., Yokohama, 244, Japan). Journal of Applied Polymer Science, 44(9), 1573-82 (English) 1992. CODEN: JAPNAB. ISSN: 0021-8995.

AB Silsesquioxane, siloxane, and silmethylene polymers with pendent phenolic groups were prepd. to obtain alkali-sol. organosilicon These polymers had structures in which the phenol polymers. moieties were sepd. by 1 carbon from the Si. The hydroxy groups were protected as methoxy groups during polymn., then were changed into hydroxy groups by a reaction with trimethylsilyl iodide

followed by alcoholysis. Silsesquioxane with phenolic hydroxy groups possessed excellent properties for matrix resins of alkali-developable organosilicon **resists**, such as 0 reactive ion etching resistance and heat resistance.

CC 35-5 (Chemistry of Synthetic High Polymers) Section cross-reference(s): 74

siloxane pendent hydroxybenzyl prepn; silsesquioxane pendent hydroxybenzyl prepn; methoxybenzyltrichlorosilane prepn polymn; polysilane pendent hydroxyphenyl prepn; pos resist organosilicon

IT Polymerization

(of methoxybenzylchlorosilanes, for hydroxybenzyl -pendent silsesquioxanes or siloxanes or polysilanes)

IT Silsesquioxanes

(hydroxybenzyl, prepn. and characterization and alkali soly. of)

IT Resists

(**pos**.-working, from organosilicons, prepn. and characterization of)

L30 ANSWER 17 OF 22 HCAPLUS COPYRIGHT 2002 ACS
1991:644060 Document No. 115:244060 Positive-working
silicon-containing photoresists. Nate, Kazuo; Mizushima,
Akiko; Sugiyama, Hisashi; Ikeda, Hiroshi (Hitachi, Ltd., Japan).
Jpn. Kokai Tokkyo Koho JP 03102354 A2 19910426 Heisei, 6 pp.
(Japanese). CODEN: JKXXAF. APPLICATION: JP 1989-239936 19890918.

AB The resists mainly contain alkali-sol. Si-contg. polymer with phenolic OH groups, alkali-sol. polymers, photo-inactivated dissoln. inhibitor, and solvent. These photoresists form thick layer without cracks, and provide high sensitivity, resoln., and resistance to 0 plasma etching. Thus, a hydrolysis product of p-methoxybenzyltrichlorosilane was treated with trimethylsilyliodide to obtain p-hydroxybenzylsilsesquioxane. A soln. contg. this silsesquioxane 80, pyrogallol resin 20, and 1,2-naphthoquinonediazide-5-sulfonate ester of 2,3,4trihydroxybenzophenone (75% triester) 30 parts was applied on Si wafer and prebaked to obtain a 20-.mu.m-thick layer. Exposure and development with 2.38% Me4NOH gave thick pattern layer without cracks, with sensitivity 600 mJ/cm2. A bilevel resist with a 5-.mu.m-thick upper layer of the invention resist was exposed with 300 mJ/cm2 and developed to obtain pattern with 5-.mu.m line-and-space. Etching in O plasma gave pattern with rectangular profile and aspect ratio .gtoreq.5.

IC ICM G03F007-022

ICS C08K005-33; C08L083-06; G03F007-075; H01L021-027

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 38

ST photoresist silicon contg crackless; silsesquioxane photoresist bilevel resist

IT Silsesquioxanes

(hydroxybenzyl, photoresists contq.,

alkali-developable, thick layer without cracks, upper layer of bilevel resists)

IT Resists

(photo-, silsesquioxane-contg., alkali-developable, thick layer without cracks, upper layer of bilevel resists)

IT 116763-64-5

(photo-decompd. dissoln. inhibitor, photoresists contg. alkali-sol. silsesquioxane and)

L30 ANSWER 18 OF 22 HCAPLUS COPYRIGHT 2002 ACS

1991:546438 Document No. 115:146438 Bilayer resist system utilizing alkali-developable organosilicon positive photoresist (OSPR). Nate, Kazuo; Mizushima, Akiko; Sugiyama, Hisashi (Prod. Eng. Res. Lab., Hitachi, Ltd., Yokohama, 244, Japan). Proceedings of SPIE-The International Society for Optical Engineering, 1466 (Adv. Resist Technol. Process. 8), 206-10 (English) 1991. CODEN: PSISDG. ISSN: 0277-786X.

AB A bilayer resist system utilizing an alkali-developable organosilicon pos. photoresist (OSPR) was developed. The composite prepd. from an alkali-sol. organosilicon polymer, poly(p-hydroxybenzylsilsesquioxane) and naphthoquinone diazide becomes an alkali-developable pos. photoresist which is sensitive to UV (i-line and g-line) region and exhibits high oxygen reactive ion etching resistance. The sensitivity and the resoln. of OSPR are almost the same as those of conventional novolak-based pos. photoresists.

A bilayer resist system utilizing OSPR as the top imaging

layer gave fine patterns of org. underlayers with high aspect ratio. 74-5 (Radiation Chemistry, Photochemistry, and Photographic and

Other Reprographic Processes)
Section cross-reference(s): 76

ST polyhydroxybenzylsilsesquioxane pos photoresist submicron lithog; organosilicon bilayer photoresist

IT Silsesquioxanes

(hydroxybenzyl, pos. bilayer photoresists compn. contg., as top imaging layer)

IT Resists

CC

(photo-, pos.-working, bilayer, contg. poly(
hydroxybenzylsilsesquioxane) as top imaging layer and
naphthoquinonediazidesulfonyl chloride-trihydroxybenzophenone
ester)

IT 5610-94-6

(pos. photoresist compn. contg. poly(hydroxybenzylsilsesquioxane) and)

L30 ANSWER 19 OF 22 HCAPLUS COPYRIGHT 2002 ACS
1990:28233 Document No. 112:28233 Excimer laser-sensitive agents.
Sugyama, Hisashi; Ehata, Keisuke; Nate, Kazuo; Mizushima, Akiko
(Hitachi, Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 01155337 A2
19890619 Heisei, 6 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP
1987-314002 19871214.

- * STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY AVAILABLE VIA OFFLINE PRINT *
- Compds. derived from satd. hydrocarbon skeletons having OH groups, AΒ that have an aliph. diazoketone group bonded to the skeleton though ester formation with the OH groups, are used as the title agents. Typically, these compds. are derived from cholic, deoxycholic, lithocholic acids or their derivs., the diazoketone groups being R1C(O)C(N2) (R1 = org. group not contg. arom. group), and the compds. are preferably I, II, or III R2 = H, C1-10 alkyl; R3 = H, RC(0)C(N2)CO (not H simultaneously) /. These agents are easily bleached by <300 nm radiation, have good solvent soly. and film-forming properties, do not sublime, and effectively inhibit dissoln. of alkali-sol resins in their presence, and therefore are conveniently used in, e.g., pos.-working resists patterned by a KrF excimer laser. Thus, I [R2 = Me; R3 = OC(O)C(N2)OMe] were highly sol. in many org. solvents, with an absorption max. at 258 nm, /and irradn. at 249 nm produced rapid bleaching. The decrease of thickness of a film of poly(phydroxybenzylsilsesquioxane) by immersion in 0.05M (HOCH2CH2) 4NOH was decreased to 1/19, by the presence of the above invention compd.

IC ICM G03C001-72 ICS G03C005-16

- CC 74-12 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- ST diazoketonic ester photosensitive UV bleachable; photoresist pos excimer laser sensitive

IT Silsesquioxanes

(hydroxybenzyl dissoln. inhibition of, in alk. soln., by UV-bleachable diazoketonic esters of cholic acid derivs.)

IT Resists

(photo-, **pos**.-working, excimer laser-sensitive, diazoketonic esters of cholic acid derivs. as)

- L30 ANSWER 20 OF 22 HCAPLUS COPYRIGHT 2002 ACS
- 1989:564040 Document No. 111:164040 Aliphatic diazoketones for KrF excimer laser lithography. Sugiyama, Hisashi; Ebata, Keisuke; Mizushima, Akiko; Nate, Kazuo (Prod. Eng. Res. Lab., Hitachi Ltd., Yokohama, 244, Japan). Journal of Photopolymer Science and Technology, 2(3), 391-400 (Japanese) 1989. CODEN: JSTEEW. ISSN: 0914-9244.
- AB A new class of sensitizers for **pos**. KrF excimer laser **resist** is described. Novel .alpha.-diazoacetoacetate derivs. of steroids and aliph. polyfunctional alcs. were synthesized. These compds. underwent photolysis upon UV exposure to give carboxylic acids and showed good bleaching properties. The concn. of photoactive groups in a sensitizer mol. should be reduced

to obtain high resist transparency and enhanced bleaching activity. The steroids showed good dissoln -inhibiting property in poly(p-hydroxybenzylsilsesquioxane) (HSQ), but not in poly(vinylphenol), that was superior to the polyfunctional alc. systems. In the steroid systems, the resists gained sensitivity with increasing no. of photoactive groups in a sensitizer mol., but leveled off when the no. of groups reached 3. Methylcholatetris(.alpha.-diazoacetoacetate) was found to be the best sensitizer for pos. KrF excimer laser resist

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

IT Diazo compounds

(diazoacetoacetate derivs. of steroids and aliph. polyfunctional alcs., as photosensitizers for photoresists for deep-UV lithog.)

IT Silsesquioxanes

(hydroxybenzyl, photoresist/for deep-UV

lithog. from, diazoacetoacetate photosensitizers for)

IT Resists

(photo-, for deep-UV lithog., diazoacetoacetate derivs. as photosensitizers for)

L30 ANSWER 21 OF 22 HCAPLUS COPYRIGHT 2002 ACS

1989:564016 Document No. 111:164016 **Positive** excimer laser **resists** prepared with aliphatic diazoketones. Sugiyama, Hisashi; Ebata, Keisuke; Mizushima, Akiko; Nate, Kazuo (Prod. Eng. Res. Lab., Hitachi, Ltd., Yokohama, 244, Japan). Polymer Engineering and Science, 29(13), 863-7 (English) 1989. CODEN: PYESAZ. ISSN: 0032-3888.

AB A new class of alkali-developed pos. excimer laser (KrF)
resists is described. Novel .alpha. diazoacetoacetates
derived from aliph. polyfunctional alcs. were synthesized. These
compds. undergo photolysis upon deep-UV exposure to yield carboxylic
acids, and exhibit excellent bleaching effects. Some of them, esp.
those having steroid skeletons, act as effective dissoln.
inhibitors. The composites prepd. from these compds. and poly(phydroxybenzylsilsesquioxane) were used as alkali-developable
pos. deep-UV resists, whose sensitivities depend
on the no. of photoactive groups in one photoactive mol. Imaging
results of KrF excimer laser projection printing are presented.

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST lithog deep UV resist diazoketone; photoactive diazoacetoacetate lithog photoresist; pos photoresist excimer laser lithog

IT Silsesquioxanes

(lithog. pos.-working excimer laser photoresist contg. hydroxybenzylsilsesquioxane matrix in)

IT Diazo compounds

(photoactive diethylacetoacetates, for deep-UV photoresist applications, lithog. characteristics of)

IT Silsesquioxanes

(hydroxybenzyl, pos. excimer laser

resists prepd. with aliph. diazoketones in)

IT Resists

(photo-, pos.-working, for excimer laser exposure, with hydroxybenzylsilsesquioxane matrix and photoactive aliph. diazoketones, lithog. characteristics of)

IT 123131-56-6 123131-57-7 123131-58-8 123131-59-9 123153-94-6 123153-95-7 123153-96-8 123153-97-9 123153-98-0 123153-99-1 123154-00-7 123154-01-8

(lithog. pos.-working excimer laser photoresist contg. hydroxybenzylsilsesquioxane matrix in)

L30 ANSWER 22 OF 22 HCAPLUS COPYRIGHT 2002 ACS

- 1989:202623 Document No. 110:202623 Alkali-developable organosilicon positive photoresist (OSPR). Sugiyama, Hisashi;
 Inoue, Takashi; Mizushima, Akiko; Nate, Kazuo (Prod. Eng. Res. Lab., Hitachi, Ltd., Yokohama, 244, Japan). Proceedings of SPIE-The International Society for Optical Engineering, 920 (Adv. Resist Technol. Process. 5), 268-73 (English) 1988. CODEN: PSISDG. ISSN: 0277-786X.
- AB A new alkali-developable organosilicon pos. photoresist for a bilayer resist system was developed. Novel alkali-sol. organosilicon polymers, polysilsesquioxane, polysiloxane, and polysilmethylene, were prepd. as the matrix polymers. Among these polymers, poly(phydroxybenzylsilsesquioxane) (I) exhibited the highest O reactive ion etching (RIE) resistance / A composite (OSPR-1334) prepd. from I and naphthoquinone diazide becomes an alkali-developable pos. photoresist which is sensitive to i - g line light. The sensitivity and the resoln. of OSPR-1334 were almost the same as those of conventional novolak-based resists when aq. tetra(2hydroxyethyl) ammonium hydroxide is used as the developer. OSPR-1334 had excellent resistance to O2RIE. The etch rate was 3.6 nm/min, while that of polyimide or novolak-based resists was 100 nm/min. Thus, OSPR-1334 is suitable for use as the top layer of a bilayer resist system. Submicron patterns with high aspect ratio can be easily obtained with this new bilayer resist system.
- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- ST photoresist organosilicon alkali developable lithog; hydroxybenzylsilsesquioxane photoresist bilayer lithog; silsesquioxane hydroxybenzyl photoresist bilayer lithog

IT Silsesquioxanes

(lithog. **photoresist**, prepn. and characterization of, for bilayer system)

IT Siloxanes and Silicones, properties ((hydroxyphenyl)methyl Me, lithog. characterization of, for resist applications) Siloxanes and Silicones, properties IT (di-Me, (hydroxyphenyl) hydrogen, lithog. characterization of, for resist applications) IT Silsesquioxanes (hydroxybenzyl, pos. photoresist alkali-developable, for bilayer lithog. resist process) IT Resists (photo-, pos.-working, alkali-developable organosilicon, contg. poly(hydrozybenzylsilsesquioxane), for application in bilayer systems) IT120366-99-6, OSPR 1334 (lithog. bilayer photoresist system with top layer of, characteristics of) 75-59-2, Tetramethylammońium hydroxide IT 123-41-1 631-41-4 33667-48-0 (lithog. developer compn/contg., for organosilicon pos . photoresists from poly(hydroxybenzylsilsesquioxane)) 1143-72-2D, 2,3,4-Trihydroxybenzophenone, reaction products with IT naphthoquinonediazide sulfonyl chloride 3770-97-6D, reaction products with trihydroxybenzophenone 5610-94-6 (photoresist from poly(hydroxybenzylsilsesquioxane

and sensitizer system contg.)